

1-ARYL- OR 1-ALKYLSULFONYL-HETEROCYCLYL BENZAZOLES AS
5-HYDROXYTRYPTAMINE-6 LIGANDS

BACKGROUND OF THE INVENTION

5 This application claims priority from copending application serial number 60/245118, filed on November 2, 2000, the entire disclosure of which is hereby incorporated by reference.

10 Compounds capable of forming 5-HT6 receptor ligands are potentially useful in the treatment of a number of central nervous system disorders such as anxiety, depression, epilepsy obsessive compulsive disorders, migraine, cognitive disorders, sleep disorders, feeding disorders, panic attacks, disorders resulting from 15 withdrawal from drug abuse, schizophrenia, or certain gastrointestinal disorders such as irritable bowel syndrome. Significant efforts are being made to understand the recently identified 5HT-6 receptor and its possible role in neuropsychiatric and neurodegenerative 20 functions. To that end, new compounds which demonstrate a binding affinity for the 5HT-6 receptor are earnestly sought, particularly as potential potent therapeutic agents.

25 Therefore, it is an object of this invention to provide compounds which are useful as therapeutic agents in the treatment of a variety of conditions related to or affected by the 5-HT6 receptor.

It is another object of this invention to provide methods and compositions useful for the treatment of

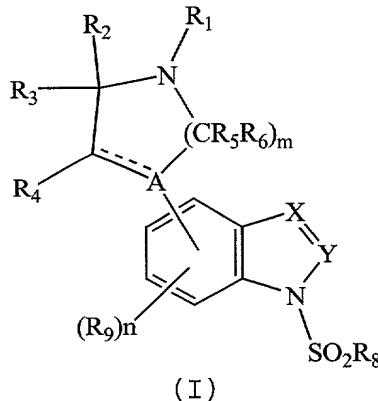
psychoses (e.g., schizophrenia, anxiety, or depression), motor disorders (e.g., Parkinson's disease), anxiety, depression, obsessive compulsive disorder, attention deficit disorder, or any condition which is known to be related to or affected by the 5-HT₆ receptor.

These and other objects and features of this invention will become more apparent by the detailed description set forth hereinbelow.

10

SUMMARY OF THE INVENTION

The present invention provides a compound of formula I



15

wherein

A is C, CR₁₀ or N;

X is CR₁₁ or N;

20 Y is CR₇ or N with the proviso that when X is N, then
Y must be CR₇;

R₁ is H, C₁-C₆alkylcarbonyl, C₁-C₆alkylcarbonyloxy or
an C₁-C₆alkyl, C₁-C₆alkenyl, C₁-C₆alkynyl or C₅-
C₇cycloheteroalkyl group each optionally
substituted;

25

R₂, R₃, R₄, R₅ and R₆ are each independently H, halogen, OH or an optionally substituted C₁-C₆alkyl group;

R₇ and R₁₁ are each independently H, halogen or an C₁-C₆alkyl, aryl, heteroaryl or C₁-C₆alkoxy group each optionally substituted;

R₈ is an C₁-C₆alkyl, aryl or heteroaryl group each optionally substituted;

R₉ is H, halogen or a C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆alkenyl, aryl or heteroaryl group each optionally substituted;

R₁₀ is H, OH or an optionally substituted alkoxy group;

m is an integer of 1, 2 or 3;

n is 0 or an integer of 1, 2 or 3; and

---- represents a single bond or a double bond; or a pharmaceutically acceptable salt thereof.

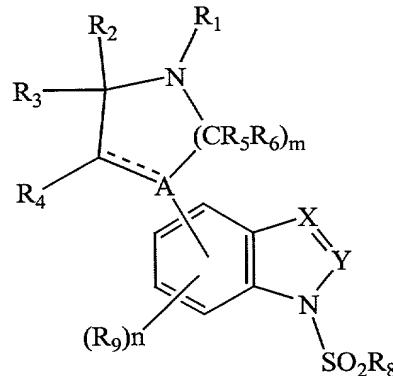
The present invention also provides methods and compositions useful in the treatment of central nervous system disorders.

DETAILED DESCRIPTION OF THE INVENTION

The 5-hydroxytryptamine-6 (5-HT6) receptor is one of the most recent receptors to be identified by molecular cloning. Its ability to bind a wide range of therapeutic compounds used in psychiatry, coupled with its intriguing distribution in the brain has stimulated significant interest in new compounds which are capable of interacting with or affecting said receptor. At present, there are no known fully selective agonists. Significant

efforts are being made to understand the possible role of the 5-HT₆ receptor in psychiatry, cognitive dysfunction, motor function and control, memory, mood and the like. To that end, compounds which demonstrate a binding 5 affinity for the 5-HT₆ receptor are earnestly sought both as an aid in the study of the 5-HT₆ receptor and as potential therapeutic agents in the treatment of central nervous system disorders.

Surprisingly, it has now been found that 1-alkyl- or 10 1-arylsulfonyl-heterocyclylbenzazoles of formula I demonstrate 5-HT₆ affinity along with significant subtype selectivity. Advantageously, said formula I benzazoles are effective therapeutic agents for the treatment of central nervous system disorders associated 15 with or affected by the 5-HT₆ receptor. Accordingly, the present invention provides 1-alkyl- or 1-arylsulfonyl-heterocyclylbenzazole compounds of formula I



20

(I)

wherein

A is C, CR₁₀ or N;X is CR₁₁ or N;

Y is CR₇ or N with the proviso that when X is N, then
Y must be CR₇;

R₁ is H, C₁-C₆alkylcarbonyl, C₁-C₆alkylcarbonyloxy or
a C₁-C₆alkyl, C₁-C₆alkenyl, C₁-C₆alkynyl or
5 cycloheteroalkyl group each optionally
substituted;

R₂, R₃, R₄, R₅ and R₆ are each independently H,
halogen, OH or an optionally substituted C₁-
C₆alkyl group;

10 R₇ and R₁₁ are each independently H, halogen or an C₁-
C₆alkyl, aryl, heteroaryl or alkoxy group each
optionally substituted;

R₈ is an C₁-C₆alkyl, aryl or heteroaryl group each
optionally substituted;

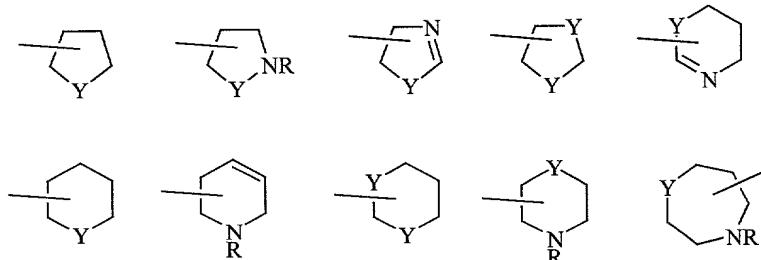
15 R₉ is H, halogen or an C₁-C₆alkyl, C₁-C₆alkoxy, C₁-
C₆alkenyl, aryl or heteroaryl group each
optionally substituted;

R₁₀ is H, OH or an optionally substituted alkoxy
group;

20 m is an integer of 1, 2 or 3;
n is 0 or an integer of 1, 2 or 3; and
---- represents a single bond or a double bond; or
a pharmaceutically acceptable salt thereof.

As used in the specification and claims, the term
25 halogen designates Br, Cl, I or F; the term aryl
designates phenyl or naphthyl; and the term
cycloheteroalkyl designates a C₅-C₇cycloalkyl ring system
containing 1 or 2 heteroatoms, which may be the same or
different, selected from N, O or S and optionally
30 containing one double bond. Exemplary of the
cycloheteroalkyl ring systems included in the term as

designated herein are the following rings wherein Y is NR, O or S and R is H or an optional substituent as described hereinbelow.



5

Similarly, as used in the specification and claims, the term heteroaryl designates a C_5 - C_{10} aromatic ring system containing 1 to 3 heteroatoms, which may be the same or different, selected from N, O or S. Such heteroaryl ring systems include pyrrolyl, azolyl, oxazolyl, thiazolyl, imidazolyl, furyl, thienyl, quinolinyl, isoquinolinyl, indolinyl, benzothienyl, benzofuranyl, benzisoxazolyl and the like; the term haloalkyl designates a C_nH_{2n+1} group having from one to $2n+1$ halogen atoms which may be the same or different; and the term haloalkoxy designates an OC_nH_{2n+1} group having from one to $2n+1$ halogen atoms which may be the same or different.

In the specification and claims, when the terms C_1 - C_6 alkyl, C_1 - C_6 alkenyl, C_1 - C_6 alkynyl, C_3 - C_7 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl are designated as being optionally substituted, the substituent groups which are optionally present may be one or more of those customarily employed in the development of pharmaceutical compounds or the modification of such compounds to

influence their structure/activity, persistence, absorption, stability or other beneficial property.

Specific examples of such substituents include halogen

atoms, nitro, cyano, thiocyanato, cyanato, hydroxyl,

5 alkyl, haloalkyl, alkoxy, haloalkoxy, amino, alkylamino, dialkylamino, formyl, alkoxycarbonyl, carboxyl, alkanoyl, alkylthio, alkylsulphinyl, alkylsulphonyl, carbamoyl, alkylamido, phenyl, phenoxy, benzyl, benzyloxy, heteroaryl, cycloheteroalkyl or cycloalkyl groups,

10 preferably halogen atoms or lower alkyl groups.

Typically, 0-3 substituents may be present. When any of the foregoing substituents represents or contains an alkyl substituent group, this may be linear or branched and may contain up to 12, preferably up to 6, more

15 preferably up to 4 carbon atoms.

Pharmaceutically acceptable salts may be any acid addition salt formed by a compound of formula I and a pharmaceutically acceptable acid such as phosphoric, sulfuric, hydrochloric, hydrobromic, citric, maleic, succinic, fumaric, acetic, lactic, nitric, sulfonic, p-toluene sulfonic, methane sulfonic acid or the like.

Preferred compounds of the invention are those compounds of formula I wherein A is N and m is 2. Also preferred are those compounds of formula I wherein R₈ is 25 an optionally substituted phenyl group and R₁ is H or a C₁-C₆alkyl or C₅-C₇cycloheteroalkyl group each optionally substituted. Further preferred compounds of the invention are those compounds of formula I wherein R₂, R₃, R₄, R₅ and R₆ are H and n is 0.

30 More preferred compounds of the invention are those compounds of formula I wherein A is N; m is 2 and R₁ is H

or a C₁-C₄alkyl or C₅-C₇cycloheteroalkyl group each optionally substituted. Another group of more preferred compounds of the invention are those compounds of formula I wherein A is N; m is 2; R₁ is H or a C₁-C₄alkyl or C₅-C₇cycloheteroalkyl group each optionally substituted; and R₈ is an optionally substituted phenyl group.

Among the preferred compounds of the invention are:

1- (phenylsulfonyl)-4-piperazin-1-yl-1H-indole;
1- [(2-bromophenyl)sulfonyl]-4-piperazin-1-yl-1H-indole;
10 1- [(6-chloroimidazo[2,1-b][1,3]thiazol-5-yl)sulfonyl]-4-
piperazin-1-yl-1H-indole;
1- [(3,4-dimethoxyphenyl)sulfonyl]-4-piperazin-1-yl-1H-
indole;
15 1- [(5-chloro-3-methyl-1-benzothien-2-yl)sulfonyl]-4-
piperazin-1-yl-1H-indole;
1- [(4-bromophenyl)sulfonyl]-4-piperazin-1-yl-1H-indole;
1- [(5-bromothien-2-yl)sulfonyl]-4-piperazin-1-yl-1H-
indole;
1- [(4,5-dichlorothien-2-yl)sulfonyl]-4-piperazin-1-yl-1H-
20 indole;
methyl 4- [(4-piperazin-1-yl-1H-indol-1-yl)sulfonyl]phenyl
ether;
4-piperazin-1-yl-1-{ [4-
25 (trifluoromethoxy)phenyl]sulfonyl }-1H-indole;
4- (4-benzylpiperazin-1-yl)-1- (phenylsulfonyl)-1H-indole;
4- (4-benzylpiperazin-1-yl)-1- [(2-bromophenyl)sulfonyl]-
1H-indole;
4- (4-benzylpiperazin-1-yl)-1- [(6-chloroimidazo[2,1-
30 b][1,3]thiazol-5-yl)sulfonyl]-1H-indole;
4- (4-benzylpiperazin-1-yl)-1- [(3,4-
dimethoxyphenyl)sulfonyl]-1H-indole;

4 - [4 - (3 - methoxybenzyl)piperazin-1-yl]-1 - (phenylsulfonyl) -
1H-indole;

1 - (phenylsulfonyl)-4 - [4 - (pyridin-4 - ylmethyl)piperazin-1 -
yl]-1H-indole;

5 1 - (phenylsulfonyl)-4 - [4 - (pyridin-3 - ylmethyl)piperazin-1 -
yl]-1H-indole;

1 - [(2 - bromophenyl)sulfonyl]-4 - [4 - (3 -
methoxybenzyl)piperazin-1-yl]-1H-indole;

1 - [(2 - bromophenyl)sulfonyl]-4 - [4 - (pyridin-4 -
ylmethyl)piperazin-1-yl]-1H-indole;

10 1 - [(2 - bromophenyl)sulfonyl]-4 - [4 - (pyridin-3 -
ylmethyl)piperazin-1-yl]-1H-indole;

1 - (phenylsulfonyl)-5-piperazin-1-yl-1H-indazole;

1 - (phenylsulfonyl)-6-piperazin-1-yl-1H-indazole;

15 1 - [(2 - bromophenyl)sulfonyl]-6-piperazin-1-yl-1H-indazole;

1 - [(4 - bromophenyl)sulfonyl]-5-piperazin-1-yl-1H-indazole;

1 - [(4 - bromophenyl)sulfonyl]-6-piperazin-1-yl-1H-indazole;

1 - [(5 - bromothien-2 - yl)sulfonyl]-5-piperazin-1-yl-1H -
indazole;

20 1 - [(5 - bromothien-2 - yl)sulfonyl]-6-piperazin-1-yl-1H -
indazole;

1 - [(4 - fluorophenyl)sulfonyl]-5-piperazin-1-yl-1H -
indazole;

1 - [(4 - fluorophenyl)sulfonyl]-6-piperazin-1-yl-1H -
indazole;

25 methyl 4 - [(5 - piperazin-1 - yl - 1H - indazol - 1 -
yl)sulfonyl]phenyl ether;

1 - phenylsulfonyl-4 - (4 - propylpiperazin-1 - yl)-1H - indazole;

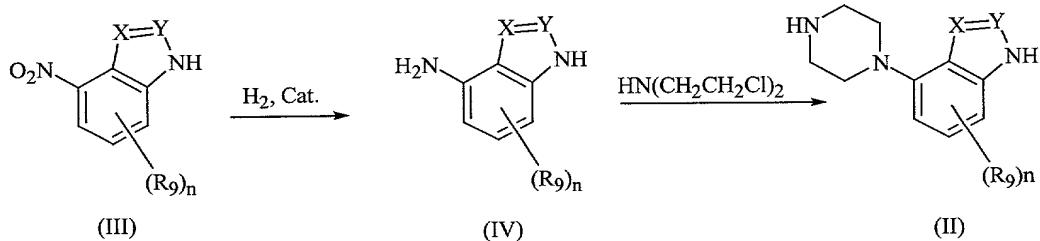
1 - phenylsulfonyl-4 - piperazin-1 - yl - 1H - indazole;

30 1 - phenylsulfonyl-4 - (4 - phenethylpiperazin-1 - yl)-1H -
indazole;

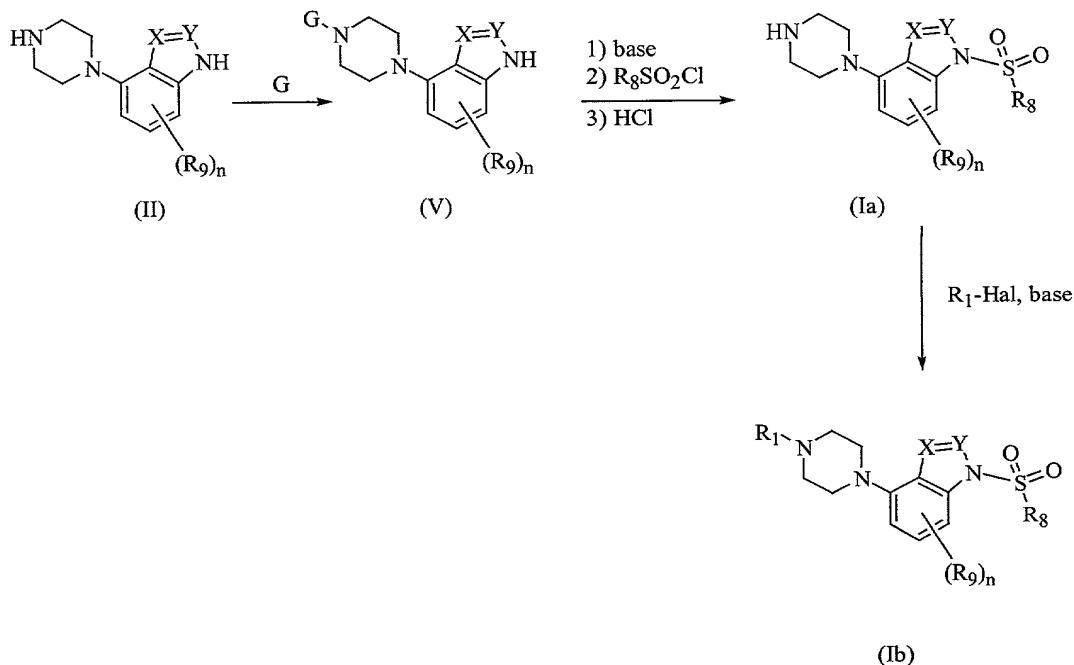
1-phenylsulfonyl-4-[4-(3-phenylpropyl)piperazin-1-yl]-1H-indazole; and

the pharmaceutically acceptable salts thereof.

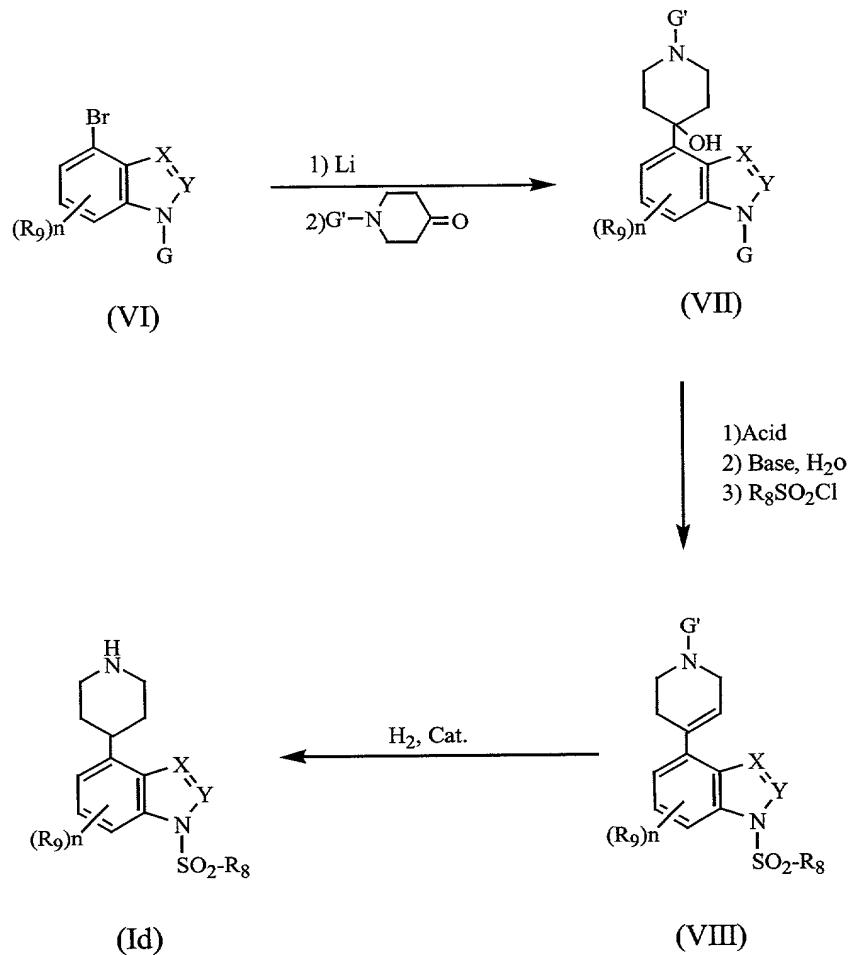
Compounds of the invention may be prepared using conventional synthetic methods and, if required, standard separation and isolation techniques. For example, 4-(piperazin-1-yl)indole compounds of formula II may be readily prepared by the catalytic hydrogenation of the 4-nitroindole precursor of formula III to the corresponding 10 4-aminoindole of formula IV and reacting said formula IV indole with a bis-alkylating agent such as bis(2-chloroethyl)amine to give the desired formula II intermediate. The reaction is illustrated in flow diagram I.

FLOW DIAGRAM I

The formula II intermediate may then be converted to a compound of formula I wherein A is N, m is 2; R₁ is H; R₂, R₃, and R₄ are H; represents a single bond; and the heterocyclyl group is in the 4-position, by reacting the formula II intermediate with a protecting group, G, for example di-t-butyl dicarbonate, to selectively protect the piperazine basic N atom to give the compound of formula V and sequentially reacting said formula V compound with a base such as NaH and a sulfonyl chloride, R₈SO₂Cl to give the protected 4-(piperazin-1-yl)-1-(substituted-sulfonyl)indole and deprotecting said indole to give the desired compound of formula Ia. Reaction of said formula Ia compound with a reagent R₁-Hal, wherein R₁ is defined hereinabove for formula I and Hal is Cl, Br or I in the presence of a base gives compounds of formula Ib wherein R₁ is other than H. The reaction sequence is shown in flow diagram II.

FLOW DIAGRAM II

5 Corresponding compounds of the invention wherein A is CR₁₀ may be obtained, for example, by lithiating a protected 4-bromoindole of formula VI wherein G is benzyl, and displacing the lithium group with a cyclic ketone such as an N-protected-4-piperidone to give the hydroxy intermediate of formula VII, which may then be dehydrated and sulfonylated in the manner described hereinabove to give the protected compound of formula VIII. Catalytic hydrogenation and simultaneous deprotection of said formula VIII compound gives the 10 desired compounds of formula I wherein _____ represents a single bond (formula Id). The reaction sequence is shown 15 in flow diagram III.

FLOW DIAGRAM III

These and other literature procedures may be
5 utilized to prepare the formula I compounds of the
invention. Employing a 5-, 6- or 7-haloindole,
-haloindazole or -halobenzimidazole substrate as starting
material and using essentially the same procedures
illustrated in flow diagrams I, II and III hereinabove
10 enables the construction of the corresponding compounds
of formula I wherein the heterocyclyl group is in the 5-,
6-, or 7-position and X or Y is N.

Advantageously, the inventive compound of formula I may be utilized in the treatment of central nervous system disorders relating to or affected by the 5-HT6 receptor such as motor, mood, psychiatric, cognitive, neurodegenerative or the like disorders. Accordingly, the present invention provides a method for the treatment of a disorder of the central nervous system (CNS) related to or affected by the 5-HT6 receptor in a patient in need thereof which comprises administering to said patient a therapeutically effective amount of a compound of formula I as described hereinabove. The compounds may be administered orally or parenterally or in any common manner known to be an effective administration of a therapeutic agent to a patient in need thereof.

The therapeutically effective amount administered in the treatment of a specific CNS disorder may vary according to the specific condition(s) being treated, the size, age and response pattern of the patient, the severity of the disorder, the judgment of the attending physician and the like. In general, effective amounts for daily oral administration may be about 0.01 to 1,000 mg/kg, preferably about 0.5 to 500 mg/kg and effective amounts for parenteral administration may be about 0.1 to 100 mg/kg, preferably about 0.5 to 50 mg/kg.

In actual practice, the compounds of the invention are administered in a solid or liquid form, either neat or in combination with one or more conventional pharmaceutical carriers or excipients. Accordingly, the present invention provides a pharmaceutical composition which comprises a pharmaceutically acceptable carrier and

an effective amount of a compound of formula I as described hereinabove.

Solid carriers suitable for use in the composition of the invention include one or more substances which may 5 also act as flavoring agents, lubricants, solubilizers, suspending agents, fillers, glidants, compression aids, binders, tablet-disintegrating agents or encapsulating materials. In powders, the carrier may be a finely divided 10 solid which is in admixture with a finely divided compound of formula I. In tablets, the formula I compound is mixed with a carrier having the necessary compression properties in suitable proportions and compacted in the shape and size desired. Said powders and tablets may contain up to 99% by weight of the 15 formula I compound. Solid carriers suitable for use in the composition of the invention include calcium phosphate, magnesium stearate, talc, sugars, lactose, dextrin, starch, gelatin, cellulose, methyl cellulose, sodium carboxymethyl cellulose, polyvinylpyrrolidone, low 20 melting waxes and ion exchange resins.

Any pharmaceutically acceptable liquid carrier suitable for preparing solutions, suspensions, emulsions, syrups and elixirs may be employed in the composition of the invention. Compounds of formula I may be dissolved 25 or suspended in a pharmaceutically acceptable liquid carrier such as water, an organic solvent, or a pharmaceutically acceptable oil or fat, or a mixture thereof. Said liquid composition may contain other suitable pharmaceutical additives such as solubilizers, 30 emulsifiers, buffers, preservatives, sweeteners, flavoring agents, suspending agents, thickening agents,

coloring agents, viscosity regulators, stabilizers, osmo-regulators, or the like. Examples of liquid carriers suitable for oral and parenteral administration include water (particularly containing additives as above, e.g.,

5 cellulose derivatives, preferably sodium carboxymethyl cellulose solution), alcohols (including monohydric alcohols and polyhydric alcohols, e.g., glycols) or their derivatives, or oils (e.g., fractionated coconut oil and arachis oil). For parenteral administration the carrier
10 may also be an oily ester such as ethyl oleate or isopropyl myristate.

Compositions of the invention which are sterile solutions or suspensions are suitable for intramuscular, intraperitoneal or subcutaneous injection. Sterile
15 solutions may also be administered intravenously.

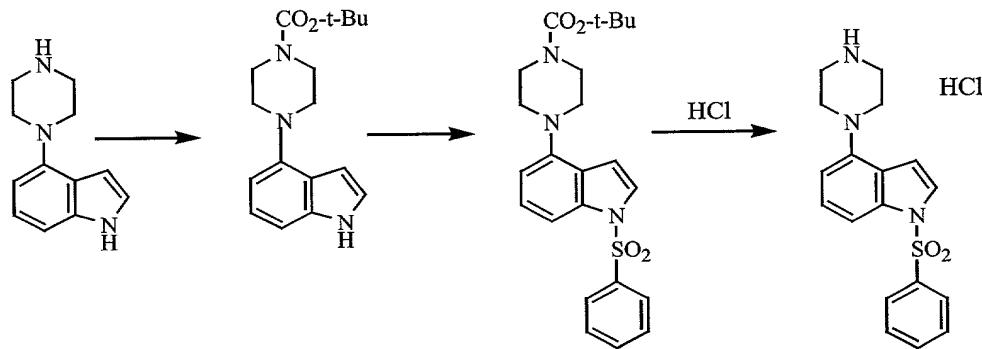
Inventive compositions suitable for oral administration may be in either liquid or solid composition form.

For a more clear understanding, and in order to illustrate the invention more clearly, specific examples
20 thereof are set forth hereinbelow. The following examples are merely illustrative and are not to be understood as limiting the scope and underlying principles of the invention in any way.

Unless otherwise stated, all parts are parts by weight. The terms HPLC and NMR designate high performance liquid chromatography and nuclear magnetic resonance, respectively.

EXAMPLE 1Preparation of 1-(Phenylsulfonyl)-4-piperazin-1-yl-1H-indole Hydrochloride

5



A mixture of 1H-indol-4-ylpiperazine (4.0 g, 20 mmol), di-t-butyl dicarbonate (4.8 g, 22 mmol) and NaOH (0.8 g, 20 mmol) in 40% dioxane is stirred at room 10 temperature for 10 hours and treated with water. The reaction mixture is extracted with ethyl acetate. The extracts are combined, dried over Na_2SO_4 and concentrated *in vacuo* to give t-butyl 4-(1H-indol-4-yl)piperazine-1-carboxylate as a colorless solid, mp 137°C, identified by 15 mass spectral and elemental analyses.

A portion of the t-butyl 4-(1H-indol-1-yl)-piperazine-1-carboxylate (1.05 g, 3.5 mmol) is added to a suspension of NaH (3.8 mmol) in tetrahydrofuran at 0°C 20 under N_2 . The resultant mixture is stirred for 0.5 hr, treated with benzenesulfonyl chloride (0.616 g, 3.5 mmol), stirred for 16 hr and treated with water. The aqueous reaction mixture is extracted with ethyl acetate. The extracts are combined, dried over Na_2SO_4 and 25 concentrated *in vacuo* to give a residue. The residue is

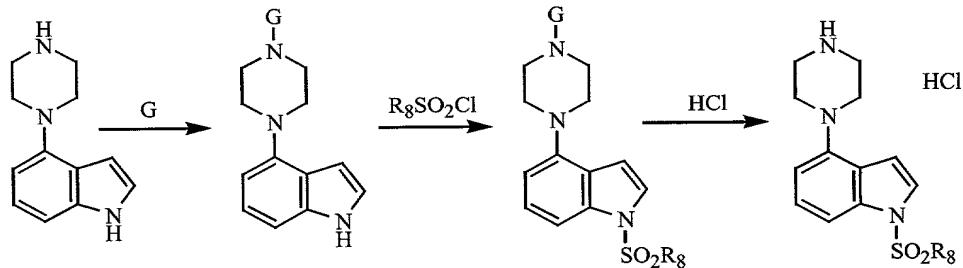
chromatographed (SiO_2 , CH_2Cl_2) to give t-butyl 4-(1-phenylsulfonyl-(1H-indol-4-yl)piperazine-1-carboxylate as a light yellow solid, 1.25 g (81% yield), mp 64-65°C, identified by mass spectral and elemental analyses.

5 A portion of the t-butyl 4-(1-benzenesulfonyl-1H-indol-4-yl)piperazine-1-carboxylate (0.85 g) is stirred in a mixture of 4N HCl and dioxane at room temperature for 2 hrs and filtered. The filtercake is dried to give the title product as a white solid, 0.64 g (99% yield) mp 10 60°C identified by mass spectral and NMR analyses.

EXAMPLES 2-13

Preparation of 1-Arylsulfonyl-4-Piperazin-1-yl)-1H-Indole

15 Hydrochloride



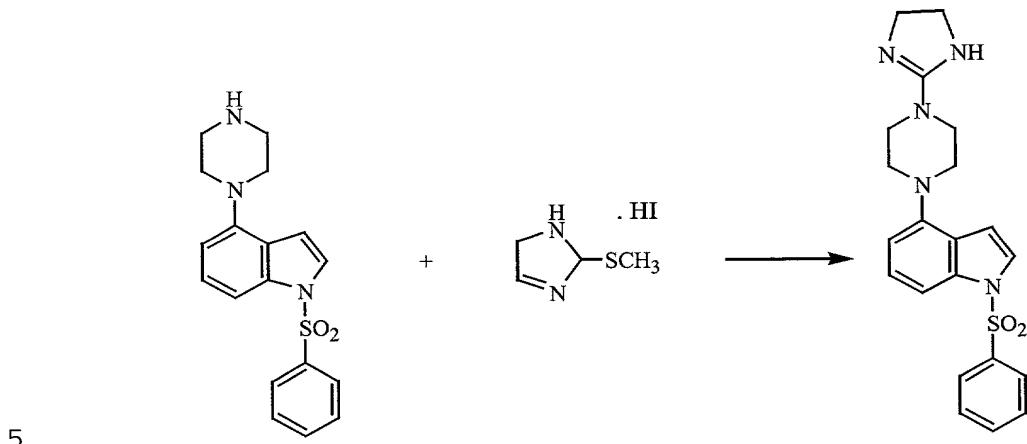
G = protecting group

Using essentially the same procedure described in Example 1 and substituting the appropriate arylsulfonyl chloride, the following compounds listed in Table I are obtained and identified by HPLC and mass spectral analyses.

TABLE I

| Ex. No. | R_8 | LCMS ¹ Min. | $M+H$ |
|------------|-------------------------------------|---------------------------|-------|
| 2 | o -bromophenyl | 2.58 | 422 |
| 3 | 6-chloroimidazo[2,1-b]thiasol-5-yl | 2.48 | 422 |
| 4 | 3,4-dimethoxyphenyl | 2.52 | 402 |
| 5 | 4-aminophenyl | 2.26 | 357 |
| 6 | benzo-2,1,3-thiazol-4-yl | | |
| 7 | benzofurazan-4-yl | | |
| 8 | 3-bromo-5-chlorothien-2-yl | | |
| 9 | 5-chloro-3-methylbenzo(b)thien-2-yl | | |
| 10 | Dansyl | | |
| 11 | 2,5-dichlorothien-3-yl | | |
| 12 | 3,5-dimethylisoxasol-4-yl | | |
| 13 | 1-methylimidazol-4-yl | | |

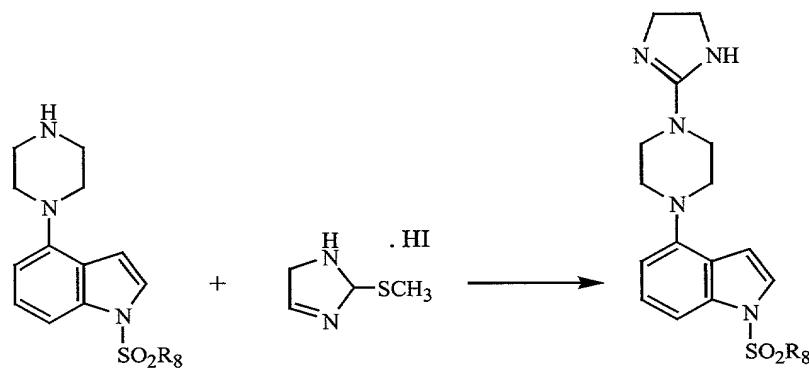
¹ LCMS conditions: Hewlett Packard 1100 MSD; YMC ODS-AM
 2.0 mm x 50 mm 5 μ column at 23°C; 3 μ L injection;
 5 Solvent A: 0.02% TFA/water; Solvent B: 0.02%
 TFA/acetonitrile; Gradient: Time 0:95% A; 0.3 min: 95%
 A; 4.7 min: 10% A, 4.9 min: 95% A; Post time 1 min.
 Flow rate 1.5 mL/min; Detection: 254 nm DAD; API-ES
 Scanning Mode Positive 150-700; Fragmentor 70 mV.

EXAMPLE 14Preparation of 4-[4-(4,5-Dihydro-1H-imidazol-2-yl)-piperazin-1-yl]-1-(phenylsulfonyl)-1H-indole

A solution of 1-(phenylsulfonyl)-4-piperazin-1-yl-1H-indole (71 mg, 0.18 mmol) in dioxane is treated with 2-methylthio-2-imidazoline hydroiodide (52.7 mg, 0.22 mmol) and N,N-diisopropylethylamine (62 μ l, 0.36 mmol), heated at 50°C for 16 hr., cooled and concentrated *in vacuo* to give a residue. The residue is purified by HPLC to give the title product, 15 mg, identified by HPLC and mass spectral analyses (2.57 min; 410 M+H) using the LCMS conditions described in Table I.

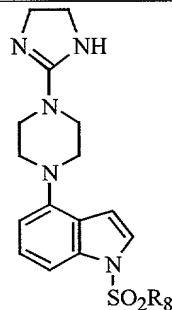
EXAMPLES 15-18Preparation of 4-Heterocyclyl-1-(arylsulfonyl)indole compounds

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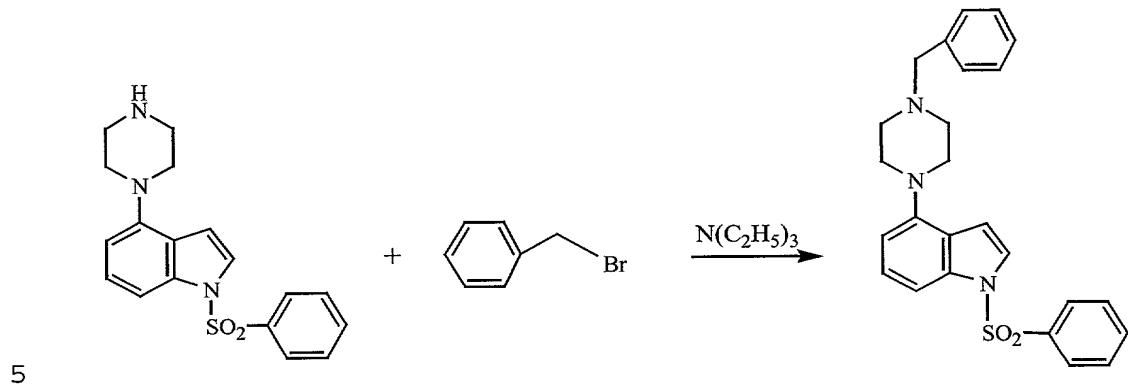
Using essentially the same procedure described in Example 14 and substituting the appropriate 1-(arylsulfonyl)indole substrate, the following compounds 10 shown in Table II are obtained and identified by HPLC and mass spectral analyses.

TABLE II



| Ex. No. | R ₈ | LCMS ¹ Min. | M+H |
|------------|------------------------------------|---------------------------|-----|
| 15 | 2-bromophenyl | 2.79 | 490 |
| 16 | 6-chloroimidazo[2,1-b]thiazol-5-yl | 2.68 | 490 |
| 17 | 3,4-dimethoxyphenyl | 2.64 | 470 |
| 18 | 4-aminophenyl | 2.46 | 425 |

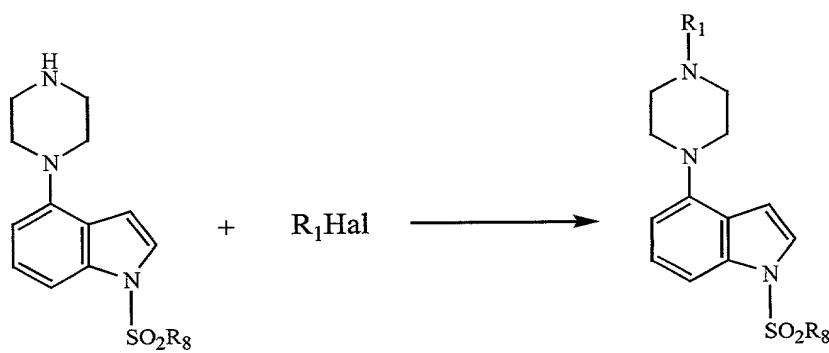
¹ LCMS conditions: same as for Table I

EXAMPLE 19Preparation of 4-(4-Benzylpiperazin-1-yl)-1-(phenylsulfonyl)-1H-indole

A solution of 1-(phenylsulfonyl)-4-piperazin-1-yl-1H-indole (71 mg, 0.18 mmol) in tetrahydrofuran is treated sequentially with benzyl bromide (21 μ l) and 10 triethyl-amine (75 μ l), shaken at room temperature for 16 hr and concentrated *in vacuo* to give a residue. The residue is purified by RP-HPLC to give the title product, 37 mg, identified by HPLC and mass spectral analyses (2.81 min; 432 M+H) using the LCMS conditions described in 15 Table I.

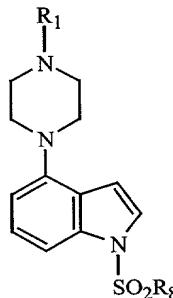
EXAMPLES 20-53Preparation of 4-Heteroaryl-1-arylsulfonylindole compounds

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Using essentially the same procedure described in Example 19 and employing the appropriate 4-(piperazin-1-yl)-1-(arylsulfonyl)indole substrate and a suitable aryl, alkyl or acyl halide, the following compounds shown in Table III are obtained and identified by HPLC and mass spectral analyses.

TABLE III

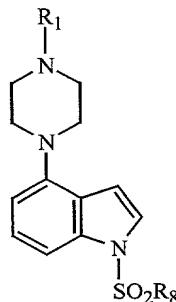


| Ex. No. | R ₁ | R ₈ | LCMS ¹ Min. M+H |
|------------|--------------------------|----------------|----------------------------------|
| 20 | 2-chloro-5-thienylmethyl | phenyl | 3.07 472 |

TABLE III (cont'd)

| Ex. No. | R ₁ | R ₈ | LCMS ¹ | |
|------------|--------------------------|-------------------------------------|-------------------|-----|
| | | | Min. | M+H |
| 21 | 3-nitrobenzyl | phenyl | 2.95 | 477 |
| 22 | Acetyl | phenyl | 3.18 | 384 |
| 23 | Benzyl | 2-bromophenyl | 2.99 | 512 |
| 24 | 2-chloro-5-thienylmethyl | 2-bromophenyl | 3.08 | 550 |
| 25 | 3-nitrobenzyl | 2-bromophenyl | 3.08 | 550 |
| 26 | Acetyl | 2-bromophenyl | 2.97 | 557 |
| 27 | Benzyl | 6-choroimidazol [2,1-b]thiazol-5-yl | 2.91 | 512 |
| 28 | 2-chloro-5-thienylmethyl | 6-choroimidazol [2,1-b]thiazol-5-yl | 3.00 | 553 |
| 29 | 3-nitrobenzyl | 6-choroimidazol [2,1-b]thiazol-5-yl | 2.87 | 557 |
| 30 | Acetyl | 6-choroimidazol [2,1-b]thiazol-5-yl | 3.23 | 464 |
| 31 | Benzyl | 3,4-dimethoxyphenyl | 2.76 | 492 |
| 32 | 2-chloro-5-thienylmethyl | 3,4-dimethoxyphenyl | 2.90 | 532 |
| 33 | 3-nitrobenzyl | 3,4-dimethoxyphenyl | 2.82 | 537 |
| 34 | Acetyl | 3,4-dimethoxyphenyl | 3.10 | 442 |

TABLE III (cont'd)

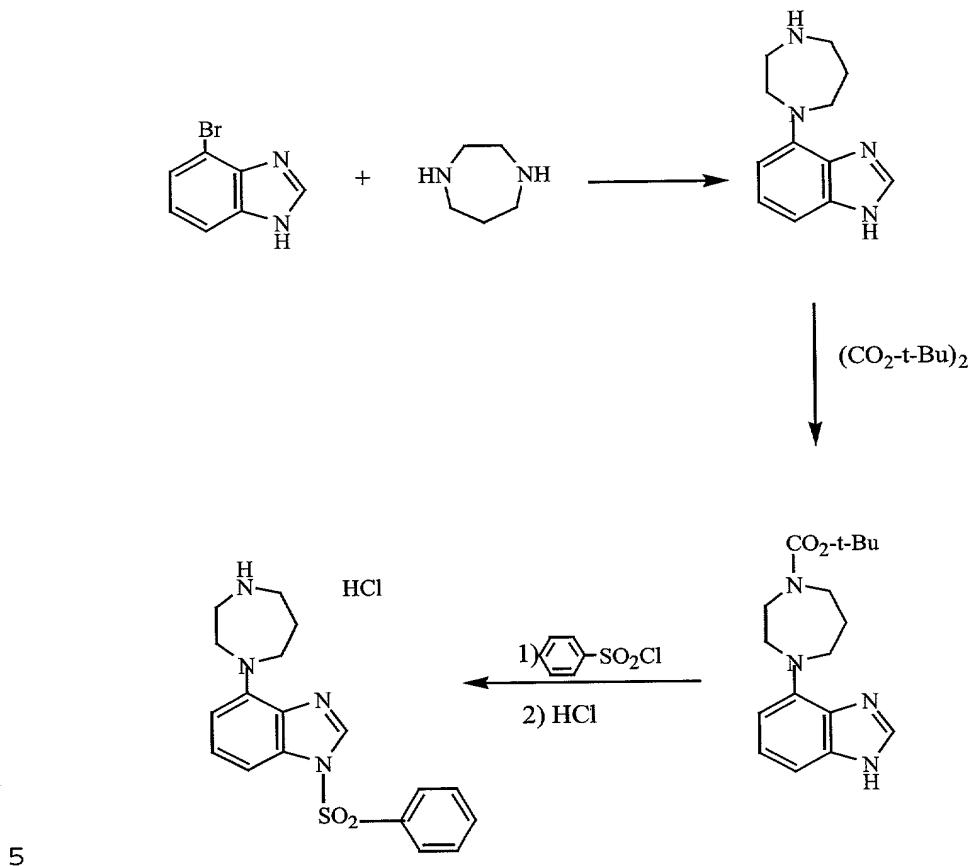


| Ex. No. | R ₁ | R ₈ | LCMS ¹ Min. | M+H |
|------------|------------------------------|--|---------------------------|-----|
| 35 | benzyl | 4-aminophenyl | 2.64 | 447 |
| 36 | methyl | 4-aminophenyl | 2.28 | 371 |
| 37 | 2-chloro-5- thienylmethyl | 4-aminophenyl | 2.82 | 487 |
| 38 | 3-nitrobenzyl | 4-aminophenyl | 2.72 | 492 |
| 39 | 3-methoxybenzyl | Phenyl | 2.88 | 462 |
| 40 | 4-pyridylmethyl | Phenyl | 2.40 | 433 |
| 41 | 3-pyridylmethyl | Phenyl | 2.42 | 433 |
| 42 | 3-methoxybenzyl | 2-bromophenyl | 2.99 | 542 |
| 43 | 4-pyridylmethyl | 2-bromophenyl | 2.51 | 513 |
| 44 | 3-pyridylmethyl | 2-bromophenyl | 2.52 | 513 |
| 45 | 3-methoxybenzyl | 6-chloroimidazo[2,1- b]thiazol-5-yl | 2.93 | 542 |
| 46 | 4-pyridylmethyl | 6-chloroimidazo[2,1- b]thiazol-5-yl | 2.48 | 513 |
| 47 | 3-pyridylmethyl | 6-chloroimidazo[2,1- b]thiazol-5-yl | 2.48 | 513 |
| 48 | 3-methoxybenzyl | 3,4-dimethoxyphenyl | 2.82 | 522 |
| 49 | 4-pyridylmethyl | 3,4-dimethoxyphenyl | 2.47 | 493 |
| 50 | 3-pyridylmethyl | 3,4-dimethoxyphenyl | 2.45 | 493 |
| 51 | 3-methoxybenzyl | 4-aminophenyl | 2.75 | 477 |

TABLE III (cont'd)

| Ex. No. | R_1 | R_8 | LCMS ¹ | |
|------------|-----------------|---------------|-------------------|-------|
| | | | Min. | $M+H$ |
| 52 | 4-pyridylmethyl | 4-aminophenyl | 2.24 | 448 |
| 53 | 3-pyridylmethyl | 4-aminophenyl | 2.26 | 448 |

¹ LCMS conditions are the same as that for Table I

EXAMPLE 54Preparation of 4-(Homopiperazin-1-yl)-1-(phenylsulfonyl)-benzimidazole hydrochloride

A suspension of 4-bromobenzimidazole (42 mmol), homopiperazine (256 mmol) and NaOt-Bu (59 mmol) in dry oxygen, under N₂, is treated with a catalytic amount of 10 Pd (OCOCH₃)₂·P(t-Bu)₃ (P/Pd = 4), heated at 120°C for 3 hr, cooled to room temperature and diluted with water. The aqueous mixture is extracted with ethyl acetate. The extracts are combined, dried over MgSO₄ and concentrated *in vacuo* to give a residue. The residue is purified by

flash chromatography to give 4-(homopiperazin-1-yl)benzimidazole.

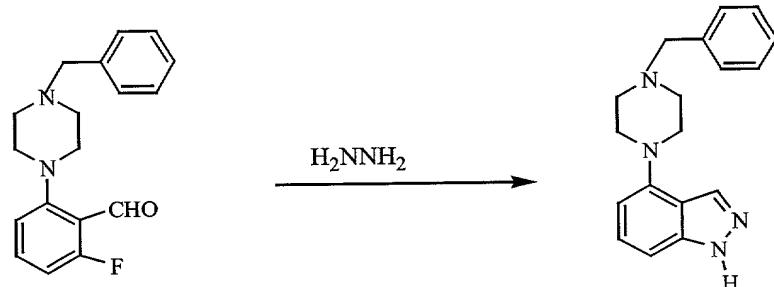
A mixture of 4-(homopiperazin-1-yl)benzimidazole (4.3 g, 20 mmol), di-t-butyl dicarbonate (4.8 g, 22 mmol) and NaOH (0.8 g, 20 mmol) in 40% aqueous dioxane is stirred at room temperature for 10 hrs and diluted with water. The aqueous mixture is extracted with ethyl acetate. The extracts are combined, dried over NaSO₄ and concentrated *in vacuo* to give t-butyl 4-(benzimidazol-4-yl)homopiperazine-1-carboxylate.

A suspension of NaH (3.8 mmol) in tetrahydrofuran at 0°C, under N₂, is treated with t-butyl 4-(benzimidazol-4-yl)-homopiperazine-1-carboxylate (1.1g, 3.5 mmol), stirred for 0.5 hr, treated with benzenesulfonyl chloride (0.616 g, 3.5 mmol), stirred for 16 hours at room temperature and diluted with water. The aqueous mixture is extracted with ethyl acetate. The extracts are combined, dried over Na₂SO₄ and concentrated *in vacuo* to give a residue. The residue is purified by flash chromatography to give t-butyl 4-(1-phenylsulfonyl)-benzimidazol-4-yl)homopiperazin-1-carboxylate.

A mixture of the thus-obtained carboxylate in 4N HCl and dioxane is stirred at room temperature for 2 hrs and filtered. The filtercake is washed with ethyl acetate and dried *in vacuo* to afford the title product.

EXAMPLE 56Preparation of 4-(4-Benzylpiperazin-1-yl)-1H-indazole

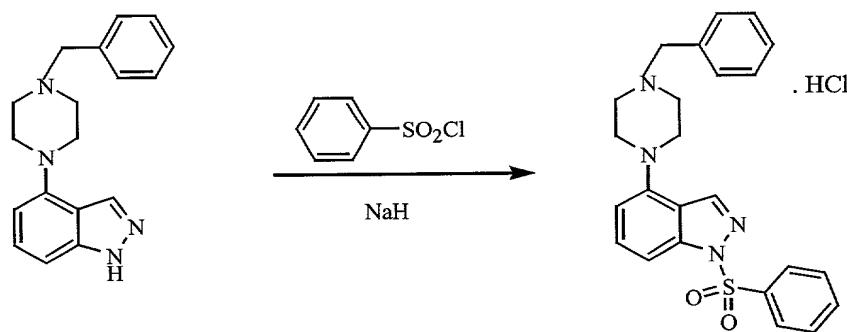
5



A stirred solution of 4-benzyl-1-(3-fluoro-2-carboxyphenyl)-piperazine (5.96 g, 20.0 mmol) in dimethylsulfoxide (10 mL) and hydrazine (10 mL) is heated at 95°C under nitrogen for 4 days. The cooled reaction is diluted with ether and washed with a mixture of water and saturated aqueous sodium bicarbonate. The organic layer is further washed sequentially with water and brine dried over MgSO₄ and concentrated *in vacuo* to give a residue. The residue is chromatographed using ethyl acetate as the eluant. The resulting oil is reconcentrated from ether to give a white foam which is stirred under hexanes/ether overnight. The resulting white powder is isolated by suction filtration and washed with hexane to give the title compound 3.11 g, (53% yield), identified by HNMR.

EXAMPLE 57Preparation of 4-(4-Benzylpiperazin-1-yl)-1-(phenylsulfonyl)-1H-indazole hydrochloride

5



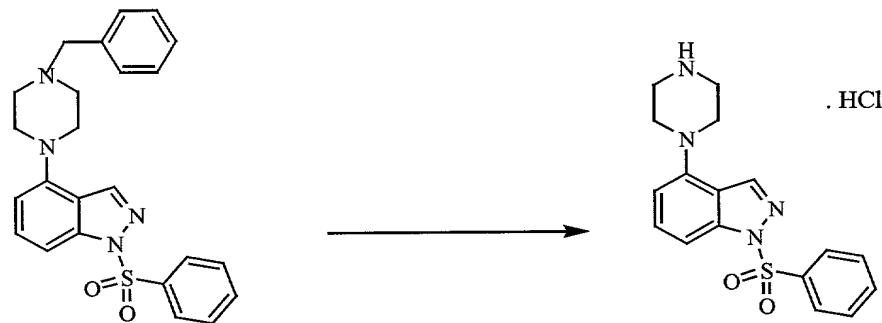
A solution of 4-(4-benzylpiperazin-1-yl)-1H-indazole (2.34 g, 8.00 mmol) in dry dimethyl formamide is treated with 0.48 g unwashed 60% NaH in mineral oil (12.0 mmol of NaH). After stirring under nitrogen for 15 min, the reaction is treated with benzenesulfonylchloride (1.53 mL, 12.0 mmol), stirred for 24 hr at ambient temperature, treated with saturated aqueous NaHCO₃ and water and extracted with ether. The organic layer is washed sequentially with water and brine, dried over MgSO₄ and concentrated *in vacuo* to give a residue. The residue is purified by flash chromatography on silica gel using 1:1 ethyl acetate:hexanes as eluant to afford the free amine of the title compound as an oil (3.14 g, 91%). A portion of this oil (432 mg, 1.0 mmol) is dissolved in ether and treated with 1.0M HCl in ether (1.1 mL, 1.1 mmol). The resulting solid is filtered, washed with ether, and dried under vacuum to provide the title compound as a light tan

solid, mp 208-209°C, identified by HNMR and mass spectral analyses.

EXAMPLE 58

5

Preparation of 1-(Phenylsulfonyl)-4-(1-piperazinyl)-1H-indazole hydrochloride

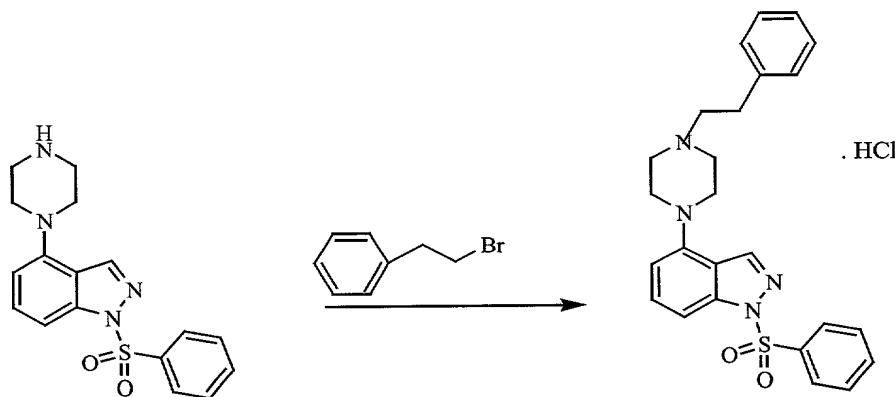


10

A solution of 1-phenylsulfonyl-4-(4-benzylpiperazin-1-yl)-1H-indazole (433 mg, 1.0 mmol) in 1,2-dichloroethane is treated with 1-chloroethyl chloroformate (0.27 mL, 2.5 mmol) heated at reflux temperature for 2 hr, and concentrated *in vacuo*. The resultant residue is heated at reflux temperature in methanol for 1.5 hr, cooled, concentrated *in vacuo* and reconcentrated from ether. The resulting tan solid is triturated with ether and crystallized from hot ethanol to give the title compound as a tan solid 237 mg (63% yield), mp 203-205 °C, identified by HNMR and mass spectral analyses.

EXAMPLE 59Preparation of 4-[4-(2-phenylethyl)piperazin-1-yl]-1-(phenylsulfonyl)-1H-indazole hydrochloride

5



A mixture of 1-phenylsulfonyl-4-piperazin-1-yl-1H-indazole (190 mg, 0.50 mmol) and K_2CO_3 (138 mg, 1.0 mmol) in dry acetonitrile is treated with phenethylbromide (0.55 mL, 2.0 mmol), heated at reflux temperature under nitrogen for 8.5 h, treated with water and extracted with methylene chloride. The combined extracts are dried over $MgSO_4$ and chromatographed on an SCX column (Varian SCX 10

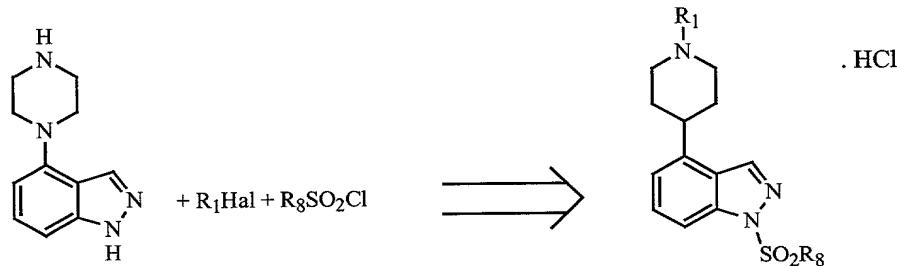
Mega Bond Elut, 5 g) eluting with ethyl acetate to remove non-basic organic material and then with 1:99 triethylamine:ethyl acetate to afford, after 15 concentration, the free amine of the title compound as a slightly yellow oil (198 mg, 89%). The oil is dissolved in ether with a small amount of ethanol to aid solubility and treated with 1.0M HCl in ether. The solution is 20 concentrated *in vacuo* and the resulting tan solid is treated with ether and suction filtered to afford the title compound as a light tan solid 209 mg, (87% yield),

mp 230-232 °C (dec), identified by NMR and mass spectral analyses.

EXAMPLES 60-72

5

Preparation of 4-Heteroaryl-1-arylsulfonylindazole compounds

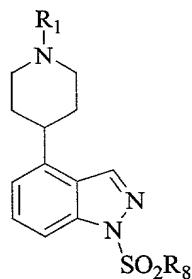


10 Using essentially the same procedures described in Examples 56-59 and employing the appropriate indazole substrate and suitable aryl, alkyl or acyl halide or arylsulfonyl chloride, the following compounds shown in Table IV are obtained and identified by NMR and mass 15 spectral analyses.

TABLE IV

| Ex. No. | R ₁ | R ₈ | mp °C | M+H |
|------------|---|----------------|----------|-----|
| 60 | 2(p-fluorophenoxy)ethyl- | Phenyl | 184-186 | 481 |
| 61 | p-fluorophenyl-CO-(CH ₂) ₃ - | Phenyl | -- | 507 |

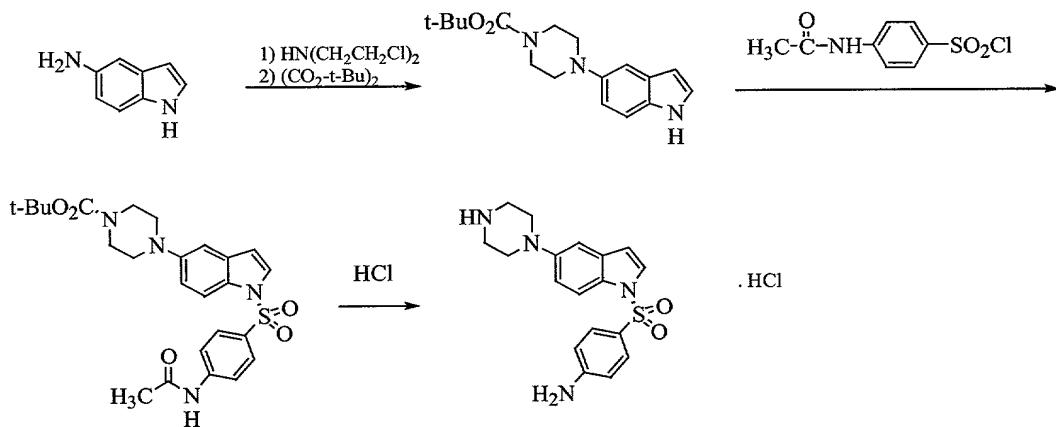
TABLE IV (cont'd)



| Ex. No. | R ₁ | R ₈ | mp °C | M+H |
|------------|-----------------------------|-----------------------------|----------|-----|
| 62 | phenyl-CO-CH ₂ - | phenyl | 202-205 | 461 |
| 63 | 3-phenylpropyl- | phenyl | 188-190 | 461 |
| 64 | n-propyl- | phenyl | 258-260 | 385 |
| 65 | benzyl | phenyl-CH=CH- | 233-235 | 459 |
| 66 | benzyl | p-fluorophenyl | 240-241 | 451 |
| 67 | benzyl | p-chlorophenyl | 238-239 | 467 |
| 68 | benzyl | naphthyl | 147-149 | 483 |
| 69 | benzyl | p-methoxyphenyl | 206-209 | 463 |
| 70 | benzyl | p-(trifluoro-methoxy)phenyl | 229-231 | 517 |
| 71 | benzyl | 2-(4,5-dichloro-thienyl)- | 235-237 | 507 |
| 72 | benzyl | p-tolyl | 215-217 | 447 |

EXAMPLE 73Preparation of 1-(4-Aminophenylsulfonyl)-5-piperazin-1-yl-1H-indole hydrochloride

5



A solution of 5-aminoindole (6.23 g, 47 mmol),
 bis(2-chloroethyl)amine hydrochloride (16.8 g, 96 mmol)
 10 and triethylamine (19 mL, 141 mmol) in butanol is heated
 at 100°C for 8 hours, cooled to room temperature and
 concentrated *in vacuo* to give 9.46 g of 5-piperazin-1-yl-
 1H-indole.

A solution of said indole in acetone and water is
 15 treated with di-*tert*-butyl dicarbonate (11.3 g, 47 mmol)
 and potassium carbonate (13 g, 96 mmol). The mixture is
 stirred at room temperature overnight, the acetone
 evaporated and the remaining aqueous phase extracted with
 ethyl acetate. The extracts are dried over MgSO₄ and
 20 concentrated *in vacuo* to give a residue. The residue is
 purified by flash chromatography to give 4-(1H-indol-5-
 yl)-piperazine-1-carboxylic acid *tert*-butyl ester.

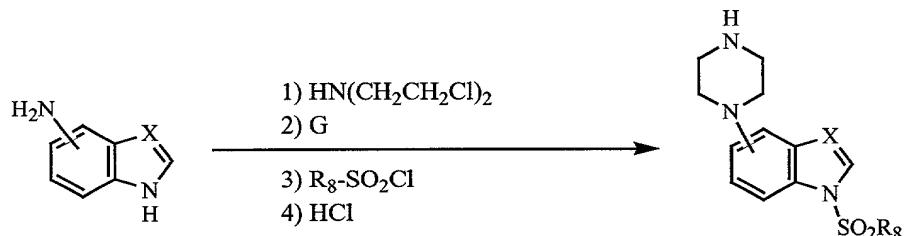
A solution of said ester (60 mg, 0.2 mmol) in tetrahydrofuran is treated with sodium hydride (30 mg, 0.5 mmol) followed by N-acetylsulfanilyl chloride (25 uL, 0.2 mmol), shaken at room temperature for 16 hours and 5 concentrated *in vacuo* to give 4-[1-(4-acetylaminophenylsulfonyl)-1H-indol-5-yl]-piperazine-1-carboxylic acid *tert*-butyl ester.

The thus-obtained ester is dissolved in methanol, treated with concentrated hydrochloric acid (100 uL), 10 shaken at 60°C for 2 hours and concentrated *in vacuo* to give a residue. The residue is purified by HPLC to give the title product, 15 mg, identified by HPLC and mass spectral analyses (r.t. 2.37 min., M+H 357).

15

EXAMPLES 74-102

Preparation of Piperazinyl-1-arylsulfonylbenzimidazole and indole compounds



G= protecting group

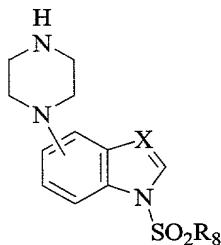
20

Using essentially the same procedures described in Example 73 and employing the appropriate aminooindole or aminobenzimidazole substrate and suitable arylsulfonylchloride reagents, the following compounds

shown in Table V are obtained and identified by HPLC and mass spectral analyses.

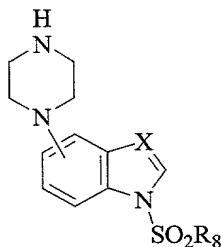
TABLE V

5



| Ex. No. | Piperazinyl | | | R8 | LCMS ¹ | |
|------------|------------------|----|--|----|-------------------|-----|
| | Ring Position | X | | | Min. | M+H |
| 74 | 5 | N | phenyl | | 1.98 | 343 |
| 75 | 6 | N | phenyl | | 1.96 | 343 |
| 76 | 5 | CH | benzo-2,1,3-thiadiazol-4-yl | | 2.56 | 400 |
| 77 | 6 | N | benzo-2,1,3-thiadiazol-4-yl | | 2.01 | 401 |
| 78 | 6 | N | 2-bromophenyl | | 2.21 | 423 |
| 79 | 5 | N | p-bromophenyl | | 2.39 | 423 |
| 80 | 6 | N | p-bromophenyl | | 2.34 | 423 |
| 81 | 5 | N | 5-bromothien-2-yl | | 2.33 | 429 |
| 82 | 6 | N | 5-bromothien-2-yl | | 2.25 | 429 |
| 83 | 5 | CH | p-(n-butoxy)phenyl | | 3.23 | 414 |
| 84 | 5 | N | p-(n-butoxy)phenyl | | 2.79 | 415 |
| 85 | 6 | N | p-(n-butoxy)phenyl | | 2.73 | 415 |
| 86 | 5 | CH | 5-chloro-1,3-dimethyl- pyrazol-4-yl | | 2.49 | 395 |
| 87 | 5 | N | 5-chloro-1,3-dimethyl- pyrazol-4-yl | | 1.88 | 396 |

TABLE V (cont'd)



| Piperazinyl | | | | LCMS ¹ | |
|-------------|------------------|----|--------------------------------------|-------------------|-----|
| Ex. No. | Ring Position | X | R ₈ | Min. | M+H |
| 88 | 5 | N | 5-chloro-3-methylbenzo-[b]thien-2-yl | 2.88 | 448 |
| 89 | 6 | N | 5-chloro-3-methylbenzo-[b]thien-2-yl | 3.10 | 448 |
| 90 | 5 | N | 2,3-dichlorothien-5-yl | 2.59 | 418 |
| 91 | 6 | N | 2,3,-dichlorothien-5-yl | 2.77 | 418 |
| 92 | 5 | N | p-fluorophenyl | 2.08 | 361 |
| 93 | 6 | N | p-fluorophenyl | 2.40 | 361 |
| 94 | 5 | N | p-methoxyphenyl | 2.11 | 373 |
| 95 | 5 | CH | 2-naphthyl | 2.92 | 392 |
| 96 | 6 | N | 2-naphthyl | 2.43 | 393 |
| 97 | 5 | CH | p-(trifluoromethoxy)phenyl | 2.97 | 426 |
| 98 | 5 | N | p-(trifluoromethoxy)phenyl | 2.57 | 427 |
| 99 | 6 | N | p-(trifluoromethoxy)phenyl | 2.54 | 427 |
| 100 | 5 | CH | p-iodophenyl | 2.92 | 468 |
| 101 | 5 | N | p-iodophenyl | 2.48 | 469 |
| 102 | 6 | N | p-iodophenyl | 2.67 | 469 |

EXAMPLE 103Comparative Evaluation of 5-HT6 Binding Affinity of Test Compounds

5

The affinity of test compounds for the serotonin 5-HT6 receptor is evaluated in the following manner.

Cultured Hela cells expressing human cloned 5-HT6 receptors are harvested and centrifuged at low speed

10 (1,000 x g) for 10.0 min to remove the culture media. The harvested cells are suspended in half volume of fresh physiological phosphate buffered saline solution and re-centrifuged at the same speed. This operation is repeated. The collected cells are then homogenized in ten
15 volumes of 50 mM Tris.HCl (pH 7.4) and 0.5 mM EDTA. The homogenate is centrifuged at 40,000 x g for 30.0 min and the precipitate is collected. The obtained pellet is resuspended in 10 volumes of Tris.HCl buffer and re-centrifuged at the same speed. The final pellet is
20 suspended in a small volume of Tris.HCl buffer and the tissue protein content is determined in aliquots of 10-25 μ l volumes. Bovine Serum Albumin is used as the standard in the protein determination according to the method described in Lowry et al., J. Biol. Chem., 193:265
25 (1951). The volume of the suspended cell membranes is adjusted to give a tissue protein concentration of 1.0 mg/ml of suspension. The prepared membrane suspension (10 times concentrated) is aliquoted in 1.0 ml volumes and stored at -70° C until used in subsequent binding
30 experiments.

Binding experiments are performed in a 96 well microtiter plate format, in a total volume of 200 μ l. To each well is added the following mixture: 80.0 μ l of incubation buffer made in 50 mM Tris.HCl buffer (pH 7.4) containing 10.0 mM MgCl₂ and 0.5 mM EDTA and 20 μ l of [³H]-LSD (S.A., 86.0 Ci/mmol, available from Amersham Life Science), 3.0 nM. The dissociation constant, K_D of the [³H]LSD at the human serotonin 5-HT₆ receptor is 2.9 nM, as determined by saturation binding with increasing concentrations of [³H]LSD. The reaction is initiated by the final addition of 100.0 μ l of tissue suspension. Nonspecific binding is measured in the presence of 10.0 μ M methiothepin. The test compounds are added in 20.0 μ l volume.

The reaction is allowed to proceed in the dark for 120 min at room temperature, at which time, the bound ligand-receptor complex is filtered off on a 96 well unifilter with a Packard Filtermate® 196 Harvester. The bound complex caught on the filter disk is allowed to air dry and the radioactivity is measured in a Packard TopCount® equipped with six photomultiplier detectors, after the addition of 40.0 μ l Microscint®-20 scintillant to each shallow well. The unifilter plate is heat-sealed and counted in a PackardTopCount® with a tritium efficiency of 31.0%.

Specific binding to the 5-HT₆ receptor is defined as the total radioactivity bound less the amount bound in the presence of 10.0 μ M unlabeled methiothepin. Binding in the presence of varying concentrations of test compound is expressed as a percentage of specific binding

in the absence of test compound. The results are plotted as log % bound versus log concentration of test compound. Nonlinear regression analysis of data points with a computer assisted program Prism® yielded both the IC₅₀ and 5 the K_i values of test compounds with 95% confidence limits. A linear regression line of data points is plotted, from which the IC₅₀ value is determined and the K_i value is determined based upon the following equation:

$$K_i = IC_{50} / (1 + L/K_D)$$

10 where L is the concentration of the radioactive ligand used and K_D is the dissociation constant of the ligand for the receptor, both expressed in nM.

15 Using this assay, the following K_i values are determined and compared to those values obtained by representative compounds known to demonstrate binding to the 5-HT₆ receptor. The data are shown in Table VI, below.

TABLE VI

| Test Compound (Ex. No.) | 5-HT ₆ binding K _i (nM) |
|----------------------------|--|
| 1 | 1.0 |
| 2 | 2.0 |
| 3 | 1.0 |
| 4 | 15.0 |
| 5 | 1.0 |
| 14 | 24.0 |
| 18 | 6.0 |

TABLE VI (cont'd)

| Test Compound (Ex. No.) | 5-HT6 binding Ki (nM) |
|----------------------------|--------------------------|
| 27 | 56.0 |
| 30 | 220.0 |
| 33 | 45.0 |
| 35 | 15.0 |
| 36 | 3.0 |
| 37 | 59.0 |
| 38 | 5.0 |
| 40 | 4.0 |
| 41 | 7.0 |
| 42 | 4.0 |
| 43 | 7.0 |
| 44 | 1.0 |
| 46 | 5.0 |
| 47 | 6.0 |
| 48 | 14.0 |
| 49 | 10.0 |
| 50 | 17.0 |
| 51 | 7.0 |
| 52 | 25.0 |
| 53 | 4.0 |
| 57 | 14 |
| 58 | 0.3 |
| 59 | 1.0 |
| 60 | 306 |
| 61 | 3.0 |

TABLE VI (cont'd)

| Test Compound (Ex. No.) | 5-HT ₆ binding Ki (nM) |
|----------------------------|--------------------------------------|
| 62 | 12 |
| 63 | 6.0 |
| 64 | 2.0 |
| 65 | 172 |
| 66 | 84 |
| 67 | 87 |
| 68 | 14 |
| 69 | 116 |
| 70 | 251 |
| 71 | 81 |
| 72 | 56 |
| 73 | 34 |
| 79 | 19 |
| 81 | 44 |
| 83 | 38 |
| 86 | 44 |
| 89 | 24 |
| 90 | 30 |
| 91 | 6 |
| 96 | 37 |
| 101 | 18 |

TABLE VI (cont'd)

| <u>Comparative Examples</u> | <u>5-HT6 binding Ki</u> |
|-----------------------------|-------------------------|
| Clozapine | 6.0 |
| Loxapine | 41.4 |
| Bromocriptine | 23.0 |
| Methiothepin | 8.3 |
| Mianserin | 44.2 |
| Olanzepine | 19.5 |

As can be seen from the results set forth above, the compounds of the present invention have a high degree of affinity for the serotonin 5-HT6 receptor sub-type.

5 Although two of the comparison compounds (clozapine and methiothepin) have similar 5-HT6 receptor affinity, they do not have the selectivity of the compounds of the present invention. The examples disclosed above 10 demonstrate up to 50-fold selectivity for the 5-HT6 receptor when compared to their affinity at the 5-HT7 receptor.